CLAIMS

1. An oral medicinal composition comprising an active ingredient and at least one foaming agent, which is ejected from a foam-developing device to be prepared into foam.

2. An oral medicinal composition according to claim 1, where the foaming agent is at least one selected from the group consisting of polyethylene glycol, saponin, sucrose esters of fatty acids, polyoxyl stearate, polyoxyethylene hydrogenated castor oil, polyoxyethylene polyoxypropylene glycol, sorbitan sesquioleate, sorbitan trioleate, sorbitan monostearate, sorbitan monopalmitate, sorbitan monolaurate, polysorbate, glyceryl monostearate, sodium lauryl sulfate and lauromacrogol.

3. An oral medicinal composition according to claim 2, where the foaming agent is at least one selected from the group consisting of polysorbate, polyethylene glycol and sodium lauryl sulfate.

4. An oral medicinal composition according to claim 3, where the foaming agent is a mixture of polyethylene glycol and polysorbate or a mixture of polyethylene glycol and sodium lauryl sulfate.

5. A method for administering an oral medicinal composition, which comprises jecting an oral medicinal composition comprising an active ingredient and at least one foaming agent from a foam-developing device to prepare the oral medicinal composition into foam for administration.

6. A method for administering an oral medicinal composition according to claim 5, where the foaming agent is at least one selected from the group consisting of polyethylene glycol, saponin, sucrose esters of fatty acids, polyoxyl stearate, polyoxyethylene hydrogenated castor oil, polyoxyethylene polyoxypropylene glycol, sorbitan sesquioleate, sorbitan trioleate, sorbitan monostearate, sorbitan monopalmitate, sorbitan monolaurate, polysorbate, glyceryl monostearate, sodium lauryl sulfate and lauromacrogol.

7. A method for administering an oral medicinal composition according to claim 6, where the foaming agent is at least one selected from the group consisting of polysorbate, polyethylene glycol and sodium lauryl sulfate.

8. A method for administering an oral medicinal composition according to claim 7, where the foaming agent is a mixture of polyethylene glycol and polysorbate or a mixture of polyethylene glycol and sodium lauryl sulfate.